

Table 2 | Critical phase 1-specific transcription factor genes in murine early T cells and their progenitors

| Gene (protein)     | Protein family            | Knockout phenotype   | Overexpression phenotype  | Selected References    |
|--------------------|---------------------------|--|---|------------------------|
| <i>Lmo2</i>        | LIM                       | • Severe stem cell defects<br>• No T cell effects if deleted after DN2 stage | T-ALL   | 1–4                    |
| <i>Gata2</i>       | GATA ZnF                  | Early stem cell defects  | Unknown   | 5,6                    |
| <i>Mef2c</i>       | MADS-box                  | T cell, B cell and NK cell defects   | • AML<br>• T-ALL association                                    | 7,8                    |
| <i>Meis1</i>       | MEIS homeodomain          | Early stem cell defects  | AML association   | 9–11                   |
| <i>Hoxa9</i>       | HOX homeobox              | • Defect in HSC proliferation<br>• Partial DN2 stage block                   | AML   | 11–14                  |
| <i>Tal1</i>        | TAL bHLH                  | Early stem cell defects  | T-ALL   | 15,16                  |
| <i>Gfi1b</i>       | • SNAG<br>• C2H2-like ZnF | HSC proliferative expansion  | Unknown   | 17                     |
| <i>Lyl1</i>        | TAL bHLH                  | Defects in LMPPs, ETPs and DN2a cells  | • B-ALL<br>• T-ALL  | 18–20                  |
| <i>Spi1</i> (PU.1) | ETS                       | • Absence of T cell and NK cell development<br>• AML                         | • Diversion to DC or myeloid cell lineages<br>• T cell lymphoma | 21–25 (reviewed in 26) |
| <i>Bcl11a</i>      | ZnF C2H2-like             | Required for B, T and NK cell development                                    | Myeloid and B cell malignancies                                 | 6,27–30                |
| <i>Hhex</i>        | Homeobox                  | Impaired monocyte development  | T-ALL   | 2,3,32                 |
| <i>Mycn</i>        | MYC bHLH                  | Defects in HSC survival and proliferation (with C-MYC)                       | AML   | 33,34                  |
| <i>Erg</i>         | ETS                       | Early stem cell defects  | T-ALL and other leukaemias                                      | 35,36                  |

AML, acute myeloid leukaemia; B-ALL, B cell acute lymphoblastic leukaemia; bHLH, basic helix–loop–helix; *Bcl11a*, B cell lymphoma–leukaemia 11A; C2H2, a form of zinc finger (Cys<sub>2</sub>His<sub>1</sub>); DC, dendritic cell; DN, double negative; ETP, early thymic progenitor; ETS, E26 transformation-specific; *Gata2*, GATA-binding protein 2; *Gfi1b*, growth factor-independent protein 1B; *Hhex*, haematopoietically expressed homeobox protein; *Hoxa9*, homeobox A9; HSC, haematopoietic stem cell; *Lmo2*, LIM domain only 2; LMPP, lymphoid-primed multipotent precursor; *Lyl1*, lymphoblastic leukaemia 1; *Mef2c*, myocyte enhancer factor 2C; *Meis1*, Meis homeobox 1; NK, natural killer; SNAG, Snail/Gfi1 domain (a repression domain); *Tal1*, gene encoding T-ALL protein 1 (also known as SCL); T-ALL, T cell acute lymphoblastic leukaemia; ZnF, zinc finger.

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